#### PATENT COOPERATION TREATY

From INTE	the RNATIONAL SEAI	RCHING AUTH	ORITY					
To: see form PCT/ISA/220				PCT  WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (PCT Rule 43bis.1)				
							Date of mailing (day/month/year) see form PCT/ISA/210 (second sheet)	
					licant's or agent's file form PCT/ISA/2			FOR FURTHER ACTION See paragraph 2 below
ł	rnational application I T/GB2004/00521		International filing date (d 08.12.2004	day/month/year)	Priority date (day/month/year) 08.12.2003	<u>-</u> 3		
International Patent Classification (IPC) or both national classification and IPC G01N33/68  Applicant OXFORD GENE TECHNOLOGY IP LIMITED								
Applicant OXFORD GENE TECHNOLOGY IP LIMITED								
This opinion contains indications relating to the following items:								
	☐ Box No. I Basis of the opinion							
	Box No. Ⅱ	Priority .				-<		
	☐ Box No. III	Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial application						
1	□ Box No. IV	Box No. IV Lack of unity of invention						
	⊠ Box No. V	Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industry applicability; citations and explanations supporting such statement				ustrial		
	☐ Box No. VI	Certain docum	nents cited					
	☐ Box No. VII	Certain defects in the international application						
	Box No. VIII Certain observations on the international application							
2.	FURTHER ACT	ION						
	written opinion o the applicant cho	vill usually be considered to be a . However, this does not apply on the chosen IPEA has notifed the national Searching Authority						
	e IPEA, the applicant is invited to nents, before the expiration of the on of 22 months from the priority	rree						
	For further options, see Form PCT/ISA/220.							
3.	For further detai	ls, see notes to	Form PCT/ISA/220.		-			
Nan	ne and mailing addre	ss of the ISA:		Authorized Officer		nes Pelag.		

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### WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/GB2004/005210

_	Box	No	o. I Basis of the opinion					
1.	With the	With regard to the language, this opinion has been established on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.						
		lan	his opinion has been established on the basis of a translation from the original language into the follow nguage , which is the language of a translation furnished for the purposes of international search under Rules 12.3 and 23.1(b)).	ring				
2.	Witl nec	re ess	egard to any <b>nucleotide and/or amino acid sequence</b> disclosed in the international application and sary to the claimed invention, this opinion has been established on the basis of:					
	a. ty	pe	pe of material:					
	[	)	a sequence listing	III B				
	(	<b>-</b>	table(s) related to the sequence listing					
b. format of material:			nat of material:	?				
	(	<b></b>	in written format	E				
	[	כ	in computer readable form	ST AVAILABLE				
	c. ti	me						
	[	כ	contained in the international application as filed.	COPY				
	{	٦.	filed together with the international application in computer readable form.	7				
	. (	]	furnished subsequently to this Authority for the purposes of search.					
3.		ha:	n addition, in the case that more than one version or copy of a sequence listing and/or table relating the as been filed or furnished, the required statements that the information in the subsequent or additional opies is identical to that in the application as filed or does not go beyond the application as filed, as ppropriate, were furnished.					
4.	Add	litio	onal comments:					
_	Во	c No	lo. II Priority					
1.	⊠	The validity of the priority claim has not been considered because the International Searching Authority does not have in its possession a copy of the earlier application whose priority has been claimed or, wher required, a translation of that earlier application. This opinion has nevertheless been established on the assumption that the relevant date (Rules 43 <i>bis.</i> 1 and 64.1) is the claimed priority date.						
2.		This opinion has been established as if no priority had been claimed due to the fact that the priority claim has been found invalid (Rules 43 <i>bis</i> .1 and 64.1). Thus for the purposes of this opinion, the international filing date indicated above is considered to be the relevant date.						
3.	Add	itio	onal observations, if necessary:					

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Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

19

lo: Claims

1-18, 20-24

Inventive step (IS)

Yes: Claims

No:

Claims

1-24

Industrial applicability (IA)

Yes: Claims

1-24

No: Claims

2. Citations and explanations

see separate sheet

## 10/581683

## AP3 Rec'd PCT/PTO 06 JUN 2005



## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (SEPARATE SHEET)

International application No.

PCT/GB2004/005210

The following documents (D) are referred to in this opinion; the numbering will be adhered to the rest of the procedure:

D1: AMERICAN SOCIETY FOR MASS SPECTROMETRY 8, 1997, PAGES 25-31

D2: RAPID COMMUNICATIONS IN MASS SPECTROMETRY 14, 2000, PAGES 924-929

D3: JOURNAL OF MASS SPECTROMETRY 37, 2002, PAGES 223-229

- 1. NOVELTY (ARTICLE 33(2) PCT)
- 1.1 Claims 1-18 and 20-24 are anticipated by D1 to D3 and are therefore not novel.

D1 (abstract; page 26, right column, last paragraph; Page 27, right column first paragraph) describes N-acetylated peptides ("label" according to claim 15) and treated with trypsin. The peptide RLAIFSC\*FR contains a deprotonated cysteic acid residue ("label", "arginine", "can form both a stabilised ion species and a protonated ion molecular species" according to claims 1, 3, 15 and 24) which balances the charge of one protonated residue, so that a further proton is incorporated without a favoured site. They were analysed using mass spectrometry ("system", "computer program" according to claims 20, 22).

D2 (abstract; Figure 2) describes the derivatisation ("label" according to claim 15) of the fibrino peptide A ("including an arginine" according to claim 15) with sulphonic acid and the sequence analysis by electro spray mass spectrometry, including single and double protonated forms ("ion species and a protonated ion molecular species that differ by one average mass unit" according to claim 15).

D3 (abstract) describes the deconvolution and deisotoping of electro spray mass spectra ("method of analysing a deisotoped peptide" according to claim 23).

1.2 The subject matter of claim 19 is not disclosed in the prior art documents and can therefore be considered as novel.



# WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (SEPARATE SHEET)

International application No.

PCT/GB2004/005210

Inventive Step (Article 33(3) PCT)

Claim 19, relating to a kit comprising a label for derivatisation, is not based on an inventive concept, since it would be obvious for the skilled person to bring together in the form of a kit, the components needed to carry out non-inventive methods. A kit for a noninventive method itself is thus not inventive.